

## § 5.25

## 21 CFR Ch. I (4–1–04 Edition)

### § 5.25 Research, investigation, and testing programs and health information and promotion programs.

(a) The following officials are authorized under sections 301, 307, 311, 1701, 1702, 1703, and 1704 of the Public Health Service Act (the PHS Act) (42 U.S.C. 241, 242l, 243, 300u, 300u-1, 300u-2, 300u-3) to establish research, investigation, and testing programs and health information and health promotion programs, which relate to their assigned functions, and to approve grants for conducting such programs:

(1) The Director, the Deputy Director for Washington Operations, and the Deputy Center Directors, Offices of Research and Management, respectively, National Center for Toxicological Research (NCTR).

(2) The Director and Deputy Directors for Science and for Regulations and Policy, Centers for Devices and Radiological Health (CDRH).

(3) The Director and Deputy Directors, Center for Biologics Evaluation and Research (CBER).

(4) The Director and Deputy Director, Center for Food Safety and Applied Nutrition (CFSAN).

(5) The Director and Deputy Director, Center for Veterinary Medicine (CVM).

(6) The Director and Deputy Director, the Directors, Office of Review Management and Office of Pharmaceutical Science, Center for Drug Evaluation and Research (CDER).

(7) The Director, Office of Orphan Products Development (OPD), Office of the Senior Associate Commissioner (OSAC), Office of the Commissioner (OC).

(b) The Director and Deputy Directors for Science and for Regulations and Policy, CDRH, are authorized to establish an electronic product radiation control program and to approve grants for conducting the program under section 532 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360ii).

(c) The Senior Associate Commissioner for Management and Systems, Office of Management and Systems (OMS), OC; the Director and Deputy Director, Office of Facilities, Acquisitions, and Central Services (OFACS), OMS, OC; the Director, Division of Contracts and Procurement Manage-

ment (DCPM), OFACS, OMS, OC; and the Chief Grants Management Officer and the Grants Management Officer, DCPM, OFACS, OMS, OC are authorized to sign and issue all notices of grant awards and amendments thereto and sign and issue notices of suspension and termination thereof for grants approved under the authority delegated in paragraphs (a) and (b) of this section.

(d) The Director, NCTR, is authorized under section 301 of the PHS Act (42 U.S.C. 241), as amended by Public Law 95-622, to make available to educational institutions, for biomedical and behavioral research, laboratory animals bred for research purposes of the Center that are not required to support Center research programs.

(e) The Senior Associate Commissioner for Management and Systems may further redelegate the authorities in paragraph (c) of this section. With the exception for paragraph (c) of this section, these officials may not further redelegate these authorities.

### § 5.26 Service fellowships.

(a) Under authority of sections 207(g) and 208(f) of the PHS Act (42 U.S.C. 209(g) and 210(f)), and within the limits of an approved service fellowship plan, the following officials are authorized to designate persons to receive service fellowships, appoint service fellows, and determine specific stipend rates for individual actions within the ranges established under an approved service fellowship plan:

(1) The Deputy Commissioner; the Senior Associate Commissioner; the Deputy Commissioner for International and Constituent Relations; the Senior Associate Commissioner for Management and Systems; the Senior Associate Commissioner for Policy, Planning, and Legislation; the Chief Counsel and Deputy Chief Counsels; and the Associate Commissioners and their Deputies.

(2) The Director, the Deputy Director for Washington Operations, the Deputy Center Directors for Research and Management, respectively, and the Associate Director, Office of Management Services, National Center for Toxicological Research (NCTR).